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Measurement of peroxides and related species in the 1993 North Atlantic Regional Experiment

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Abstract. Gaseous peroxides were measured during the North Atlantic Regional Experiment (NARE) aboard the Department of Energy's Gulfstream G-1 aircraft in August 1993. Flights originated in Halifax, Nova Scotia and covered a region extending several hundred kilometers from the eastern edge of the North American continent. Total peroxide concentration was dominated by H_2O_2. Median H_2O_2 concentration for all flights was 2.5 parts per billion by volume (ppbv), with the highest concentrations observed in spatially extensive layers between 250 and 1750 m above sea level. In the clean free troposphere, H_2O_2 concentration correlates strongly with the product O_3*H_2O, consistent with our understanding of the formation mechanism. The median peroxide concentration was 4 ppbv in pollutant plumes, with excursions above 11 ppbv. The high concentration of H_2O_2 observed, and its covariance with concentrations of anthropogenic species such as O_3, NO_y, and aerosol particles is attributed to the aging of polluted air masses advected from the continent in stable layers.

Introduction

Peroxides play an important role in atmospheric chemistry, both as oxidants for atmospheric SO_2 [*Penkett et al.*,1979; *Calvert et al.*, 1985] and as end products for the free radicals that participate in the formation of tropospheric ozone [*Kleinman*, 1986; *Madronich and Calvert*, 1990]. Although ambient peroxide concentrations have been measured in a number of field programs over the last decade, questions remain about sampling line losses, speciation of individual peroxides, and the importance of dry deposition in interpreting H_2O_2 concentrations measured at the surface.

Pre-1989 field measurements of peroxides in air and precipitation have been summarized in two review papers [*Gunz and Hoffmann*, 1990; *Sakugawa et al.*, 1990]. Peroxide concentrations at the surface correlate roughly with insolation and temperature, with deposition leading to substantial decreases at night. Many features complicate the interpretation of surface measurements, including fog and rain events, local emissions, and deposition. Peroxide measurements at high altitude surface sites, while not affected by major sources, are also complicated by cloud events and boundary layer dynamics [*Mohnen and Kadlecek*, 1989].

Heikes et al. [1987] measured gaseous peroxide concentrations up to 4 parts per billion by volume (ppbv) over the northeastern United States during fall 1984. Air near the surface had lower concentrations than air aloft, and vertical profiles frequently exhibited maximum H_2O_2 concentrations above cloud top. Lower concentrations (generally <1 ppbv) observed over the Carolinas in the winter of 1986 were attributed to lower rates of radical production [*Barth et al.*, 1989]. H_2O_2 concentrations up to 7 ppbv were observed by *P. H. Daum et al.* over Ohio during the summer of 1987. These measurements revealed a pronounced altitude dependence in peroxide concentrations with lower concentrations in the boundary layer (BL) than the free troposphere, a maximum near the top of the BL, and evidence for loss of peroxide by cloud processes. The "bulge" in peroxide concentration near the top of the BL was also observed during the same time period in the same vicinity by *Boatman et al.* [1990], and was attributed to loss by dry deposition of peroxide, reaction of its precursors with NO at lower altitudes, and lower radical production rates at higher altitudes due to decreasing water vapor concentration. Measurements in the same region during the Acid Modes study in the summer of 1988 [*Tremmel et al.*, 1993] also showed pronounced maxima in H_2O_2 concentrations at the top of the BL.

Although the aircraft measurements summarized above were carried out with a dual-channel peroxide analyzer capable of distinguishing H_2O_2 from total organic peroxides (ROOH), most report only H_2O_2 data. Exceptions include the measurements of *Barth et al.* [1989], which showed no evidence of ROOH above the detection limit of 0.1 ppbv in winter, and of *Heikes et al.* [1987] in the fall over the northeastern United States where ROOH concentrations as large as 0.5 ppbv were reported, but were not considered to be quantitative.

The August 1993 North Atlantic Regional Experiment (NARE) was an attempt to learn more about the transport and photochemistry of ozone and ozone precursors by combining observations from measurement platforms on the ground, aloft,

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and at sea. The present paper describes peroxide data obtained aboard the Department of Energy's Gulfstream G-1 aircraft, based in Halifax, Nova Scotia. A companion paper in this issue describes more fully the aircraft capabilities and instrumentation on the G-1 [*Daum et al.*, and references therein]. Synoptic patterns encountered during the mission are discussed by *Berkowitz et al.* [this issue] and *Merrill and Moody* [this issue]. A distinctive feature of the atmosphere over the western North Atlantic during this program was the frequent presence of a low-level inversion trapping warm air masses advected from the continent in isolated layers above the surface. The absence of substantial convective mixing allowed photochemical products to build up to high concentrations in these layers and preserved the relationship between peroxides and related species which would otherwise be lost through dry deposition.

Experimental

Flights

Flights of the G-1 included several west and southwesterly paths over the Bay of Fundy and Cape Sable, and several east and southeasterly paths terminating at Sable Island, approximately 250 km southeast of Halifax. Flight plans included both vertical profiles and horizontal transects, and were generally conducted in clear air, with 85% of the data collected between the hours of 1400 and 2000 UT. An overview of the G-1 flights is given in the article by *Fehsenfeld et al.* [this issue (b)].

Instrument Description

Peroxides were determined in a two-channel continuous flow analyzer of the design of G. L. Kok and A. L. Lazrus [*Kok et al.*, 1986; *Lazrus et al.*, 1986], modified to permit analysis using Fenton chemistry [*Lee et al.*, 1990; 1994] in addition to the standard p-hydroxyphenylacetic acid/peroxidase reagent in order to speciate peroxides.

A three-eighths-inch ID Teflon line, 0.75 m in length, conducted ram air to the instrument through a forward-facing inlet in the side of the plane. The instrument employs dual glass coil scrubbers to transfer gaseous peroxides into the aqueous phase, with subsequent continuous-flow derivatization to produce fluorescent products. Baselines were established several times during each flight by directing air through a Pd / alumina cartridge. Cartridge integrity was checked regularly by placing a second such cartridge upstream of the first. Signal intensity was calibrated using aqueous peroxide standards prepared daily by serial dilution of unstabilized 3% stock H_2O_2; scrubbing reagent was used for the final dilution. Stock 3% peroxide was titrated against standardized permanganate before and after the measurement period. Liquid and air flow rates were calibrated several times during the measurement period. The 0 to 95% response time of the peroxide analyzer was 70 s.

The detection limit for each channel, evaluated as three times the noise level of a zero air baseline on the ground (i.e., 3σ), was 75 parts per trillion by volume (pptv). Aloft, this value increased to 0.2 ppbv. The determination of methyl hydroperoxide (MHP) from the difference between channels had a detection limit of 0.3 ppbv. The composite error in determining the concentration of peroxide standards and liquid and air flow rates was 3.5%.

SO_2 interference was evaluated by monitoring the signal from an aqueous H_2O_2 standard in the presence of gas phase SO_2 of known concentration. No change in response was observed for SO_2 concentrations below 150 ppbv, well above the SO_2 concentrations encountered in the region.

Speciation of Peroxides

Analysis of gaseous peroxides in urban environments by the recently developed HPLC method has shown only H_2O_2, methyl hydroperoxide (CH_3OOH or MHP) and hydroxymethyl hydroperoxide ($HOCH_2OOH$ or HMHP) to be present in significant concentrations in ambient air [*Hellpointner and Gäb*, 1989; *Hewitt and Kok*, 1991; *Fels and Junkermann*, 1994]. A continuous-flow instrument capable of analyzing for these three species in real time has been developed [*Lee et al.*, 1990; 1994] but is not yet available for aircraft deployment. Therefore we analyzed for two of the three peroxides using the standard dual channel instrument. Two factors guided our selection of the two peroxides to monitor: (1) HMHP is believed to be a product of the ozonolysis of alkenes [*Simonaitis et al.*, 1991; *Hewitt and Kok*, 1991], and its concentration is expected to be greatest near the surface [*Enders et al.*, 1992]; and (2) ground measurements at the start of the program at Yarmouth, Nova Scotia indicated negligible concentrations of HMHP [J. B. *Weinstein-Lloyd et al.*, Peroxide Measurements in Yarmouth, NS, during the 1993 NARE summer intensive, manuscript to be submitted to *J. Geophys. Res.*, 1996]. Therefore the dual-channel instrument was configured to yield measurements of H_2O_2 and MHP.

Reagents employed in the dual channel system are summarized in Table 1. In channel A, the horseradish peroxidase-catalyzed reaction between peroxide and p-hydroxyphenylacetic acid (pOHPAA/HRP) produces a fluorescent dimer; and signal intensity is increased by raising pH. In Channel B, Fenton reagent, consisting of a mixture of Fe(II) and benzoic acid (FeBA) produces fluorescent hydroxy-

Table 1. Reagents for Peroxide Analyzer

CH	Scrubbing Solution*	Fluorescence Reagent*	Enhancing Reagent*	Species Detected
A	1 mM phosphate buffer, pH 9; 1 mM HCHO	3.5 mM pOHPAA; HRP 2 units/mL; pH 8.5 TRIS buffer	0.05 N NaOH	H_2O_2 + MHP
B	1 mM phosphate buffer, pH 9; 1 mM HCHO	0.8 mM $FeSO_4$; 2 mM benzoic acid; pH 1.9 H_2SO_4	6 mM $Al(NO_3)_3$; 10 mM acetate buffer; pH 3.8	H_2O_2

* all solutions prepared with conductivity grade water

benzoic acid upon reaction with H_2O_2; fluorescence intensity is enhanced in this channel by Al(III) complexation. HMHP rapidly hydrolyzes to produce H_2O_2 in the high pH scrubbing solution in both channels, causing the aqueous stream meeting fluorescence reagents to be a solution of H_2O_2 and MHP only [*Marklund*, 1971]. pOHPAA/HRP reacts with all peroxides with equal efficiency, while FeBA is specific for H_2O_2. Therefore channel A provides a measure of total peroxide, and channel B provides only H_2O_2. MHP is reported in this work as the difference between channels A and B after correcting for the collection efficiency of MHP, taking into account a recent correction to its published Henry's law constant [*Lind and Kok*, 1994; *Lind et al.*, 1986]. Two caveats should be noted when viewing the peroxide data. As discussed above, measurements at the program's start indicated negligible surface concentrations of HMHP, and we report H_2O_2 as the signal from channel B in this work. However, surface measurements may not be representative of conditions aloft, and as ambient HMHP (or other soluble, hydrolyzable peroxides) would have been collected and converted to H_2O_2 in both channels, concentrations of H_2O_2 reported here should be viewed as an upper limit. In addition, any water soluble higher peroxide that did not hydrolyze, would produce a fluorescent dimer in the pOHPAA channel, but would not produce OHBA in the FeBA channel [*Kolthoff and Medalia*, 1951]. Thus MHP concentrations reported here actually represent the sum of all nonhydrolyzable peroxides collected.

To demonstrate that MHP does not give a signal in the Fenton channel, we analyzed mixtures of H_2O_2 and MHP produced by cobalt-60 γ-radiolysis of aqueous solutions of CH_3I [*Shankar et al.*, 1969]. Irradiated samples were analyzed simultaneously in the dual-channel instrument and by high performance liquid chromatography with post-column pOHPAA/HRP derivatization; the percentage MHP determined agreed to within 15%.

Results and Discussion

Peroxides are formed by the photochemical mechanism summarized in reactions (1) - (12) below.

Initiation

$$O_3 \xrightarrow{h\nu} O_2 + O(^1D) \quad (1)$$

$$O(^1D) + H_2O \rightarrow 2\,OH \quad (2)$$

$$CH_2O \xrightarrow{h\nu} H + HCO \quad (3)$$

Formation of HO_2 Radicals

$$H + O_2 \rightarrow HO_2 \quad (4)$$

$$OH + CO \xrightarrow{O_2} HO_2 + CO_2 \quad (5)$$

Formation of RO_2 Radicals

$$OH + CH_4 \rightarrow H_2O + CH_3 \quad (6)$$

$$CH_3 + O_2 \rightarrow CH_3O_2 \quad (7)$$

Reaction with NO

$$HO_2 + NO \rightarrow OH + NO_2 \quad (8)$$

$$RO_2 + NO \rightarrow OR + NO_2 \quad (9)$$

Radical Termination Reactions

$$2\,HO_2 \rightarrow H_2O_2 + O_2 \quad (10)$$

$$CH_3O_2 + HO_2 \rightarrow CH_3OOH + O_2 \quad (11)$$

$$OH + NO_2 \rightarrow HNO_3 \quad (12)$$

H_2O_2 is formed principally by the disproportionation of HO_2 radicals, reaction (10). In polluted environments, reaction of HO_2 with NO, reaction (8), competes with disproportionation, and the subsequent photolysis of NO_2 is a major source of tropospheric ozone. The abundance of atmospheric CH_4 implies that methylperoxyl radicals (CH_3O_2) will be produced by reactions (1) and (2), followed by (6) and (7). Reaction (11) should then produce measurable quantities of MHP, which together with other peroxides can oxidize SO_2 and increase the acidity of cloud water [Lind et al., 1987]. The mechanism summarized above suggests that the eventual sink for photochemically produced free radicals must be either peroxide formation or NO_x oxidation products such as HNO_3 and peroxyacetyl nitrate (PAN).

General Observations

Concentrations of hydrogen peroxide varied from the detection limit to over 11 ppbv, with the highest concentrations observed in transported plumes of anthropogenic pollution at altitudes near 1 km. These plumes, containing high concentrations of O_3, NO_y, and related species began to be observed in the region after August 20. The early portion of the field program was characterized by light winds and anticyclonic flow, with background concentrations of photochemically related species.

A histogram of the complete data set, shown in Figure 1, exhibits a median H_2O_2 concentration of 2.5 ppbv. Median concentrations were higher in the boundary layer (2.7 ppbv) than in the free troposphere (2.2 ppbv), where boundary layer air is defined as air with dew point greater than zero. The composite vertical distribution of H_2O_2, illustrated in Figure 2, shows median peroxide concentrations between 1 and 3 ppbv. The gradual decrease in peroxide concentration with increasing altitude above the boundary layer presumably arises from decreasing OH production consistent with decreasing water vapor concentration. Deposition may contribute to the slight decrease in median concentrations observed near the surface, but episodes of high H_2O_2 concentration are evident at altitudes between 250 and 1750 m. Simultaneous measurements of other

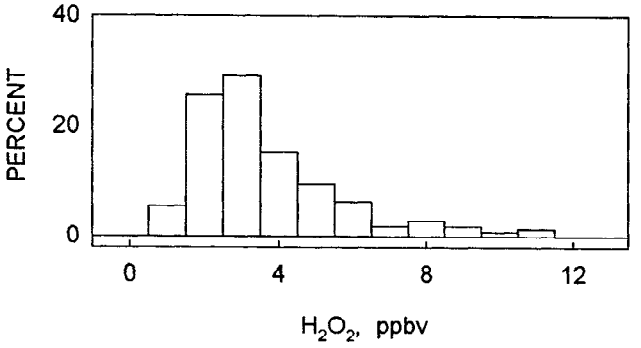

Figure 1. Frequency distribution of H_2O_2 measurements, all data.

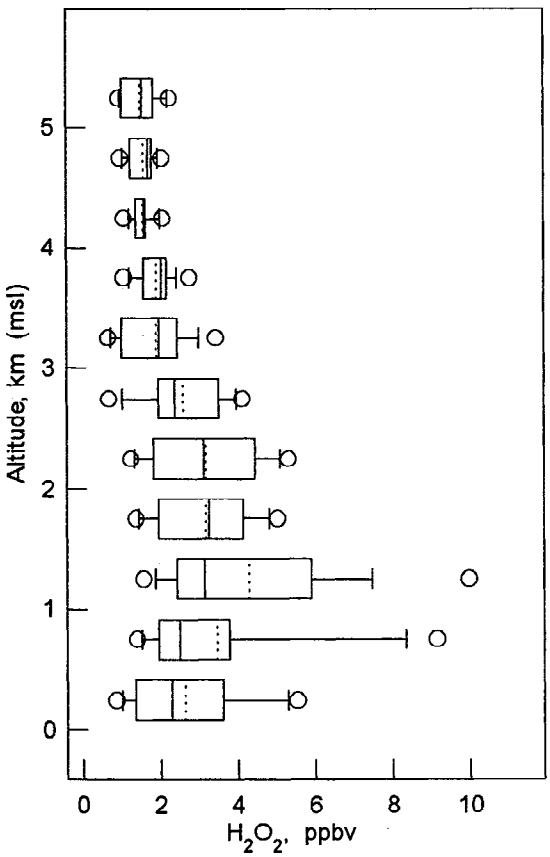

Figure 2. Composite vertical profile of peroxide measurements for the entire program. Box encloses upper and lower quartiles, with the median and mean displayed as solid and dashed lines, respectively. Extension lines enclose the 10th and 90th percentiles and circles represent outliers.

trace gas species suggest that these layers of high, variable H_2O_2 concentration are associated with plumes of anthropogenic origin [see *Daum et al.*, this issue].

MHP was detected on only 3 of the 14 flights, on August 17, 20 and 31; median concentrations of 0.45, 0.73, and 0.54 ppbv, respectively, were observed. No trends were apparent in the MHP data.

Individual Flights

Prior to August 20, flights sampled generally clean background air. On August 24, southwesterly winds began to move into the region, but it was several days before well-defined flow brought significant concentrations of pollutants from the continent. Several plumes were observed on flights from August 25 through August 31. These air masses were characterized by high concentrations of particles and NO_y, and most were well aged, as evidenced by the high levels of ozone and the large fractional conversion of NO_x to HNO_3. These regions also exhibited high peroxide concentrations. The photochemical mechanism for peroxide formation suggests that fresh air masses with high NO_x should be characterized by high O_3 from NO_2 photolysis and by low H_2O_2 concentration because of the competition for HO_2 radicals. The inverse relationship between H_2O_2 and NO_y and between H_2O_2 and O_3 is not expected to persist after the air mass leaves the source region as subsequent photochemistry produces additional H_2O_2 in NO_x-depleted air

Our measurements show a variety of relationships between H_2O_2, NO_y, and O_3 in polluted air masses. Several case studies are examined below to illustrate the range of behavior observed.

August 18. Background air was sampled on the August 18 flight between Halifax and Sable Island. This flight was typical of the early portion of the measurement program, with concentrations of NO_y below 1 ppbv, and accumulation mode aerosol particle number concentration (measured by PMS ASASP) generally below 200 cm^{-3}. A layer of very dry air (dew point -20° to -30°C) was traversed on the forward leg at an altitude of 1500 m and during the return leg at 2150 m (regions A and B in Figure 3). Despite ozone concentrations in excess of 50 ppbv in this region, H_2O_2 remained below 2 ppbv, presumably due to low water vapor concentration

The 2150-m transect crossed a sharp boundary between air masses at 1800 UT (point C in Figure 3), where dew point changed from -30° to 0°C, corresponding to a 20-fold increase in water vapor mixing ratio. At the same time, H_2O_2 concentration increased by a factor of 3, from 1 to 3 ppbv, and ozone concentration dropped from 57 to 32 ppbv. The high-ozone air mass may have descended from higher altitudes with peroxide formation inhibited by dryness. Alternatively, it may be photochemically aged air that has lost H_2O_2 and NO_x oxidation products through wet deposition.

August 27. The August 27 flight consisted of a profile over Halifax, followed by a flight to the southwest at an altitude of 1050 m to a point 150 km due east of Boston, another profile to 3800 m, and a return to the southern tip of Nova Scotia at 145 m. The initial flight leg (region A of Figure 4) was characterized by concentrations of O_3 around 50 ppbv, H_2O_2 around 5 ppbv and NO_y below 2 ppbv. A low-lying layer of pollutants was encountered during a descent to an altitude of 145 m and the return flight (region B in Figure 4) was made in this layer. Ozone concentration exceeded 100 ppbv, NO_y reached nearly 20 ppbv, and H_2O_2 concentration decreased to approximately 1 ppbv. The highly layered structure, evident in the much lower concentrations of NO_y and O_3 on the forward transect, is illustrated in Figure 5 by the vertical distributions of H_2O_2, O_3 and NO_y obtained during a sounding at 42°N, 69°W, point C in Figure 4.

Figure 6 shows that the concentration of H_2O_2 varied inversely with that of NO_x ($r^2 = 0.97$) during the flight through the low-lying layer, as one would expect from the competition between reactions *(8)* and *(10)* if this were a fresh air mass. H_2O_2 was also strongly anticorrelated with O_3 in the plume ($r^2 = 0.93$), consistent with the production of O_3 from NO_2 photolysis. Back trajectories place this air mass over the continent only 6 hours prior to our measurements, and NO_x contributes nearly 2 ppbv to a peak NO_y concentration of 16 ppbv.

August 28. Elevated concentrations of pollutants were also evident on August 28 on a flight south and east of Halifax. Concentrations of H_2O_2 and related species are illustrated for the entire flight in Figure 7. The flight path included a southbound leg at an altitude of 630 m and a return at 1240 m, with two soundings to probe the vertical boundaries of the layer. The second of these profiles, conducted at point A in Figure 7, passed through a moist air mass trapped between two inversions. Figure 8 shows that the layer is ~1 km thick, lying between 500 and 1500 m. In the trapped layer, peak concentrations of H_2O_2 and NO_y reached 9 ppbv, and O_3 reached 120 ppbv. Even higher concentrations were observed during the 630 m transect (region B of Figure 7), with ozone peaking at ~145, NO_y at ~17,

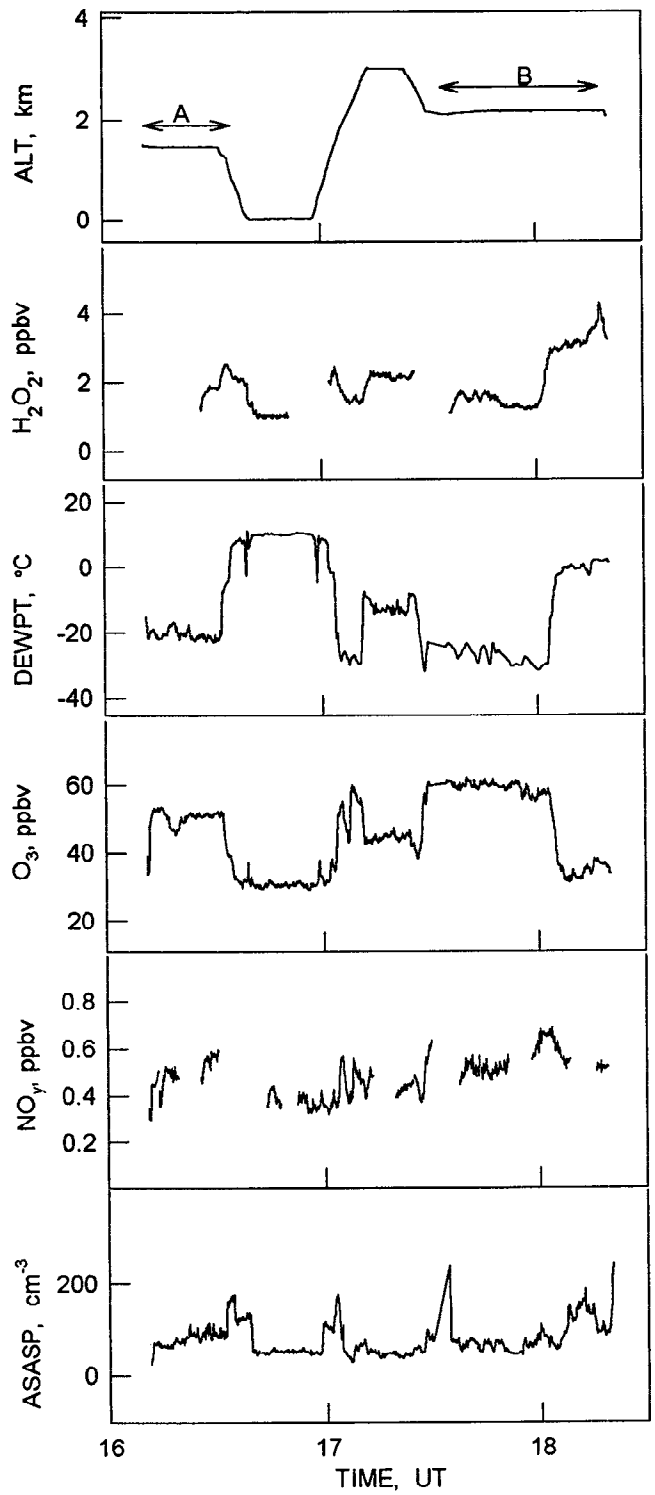

Figure 3. Concentrations of selected species on August 18 as a function of time. Horizontal transects, indicated as regions A and B, are described in the text.

and H_2O_2 at ~10 ppbv. Pollutant concentrations were also elevated during the return trip at 1240 m (region C in Figure 7), though not as high as on the lower altitude portion. A horizontal boundary was crossed during a short eastward leg at the same altitude (point D of Figure 4), with an abrupt decrease of roughly 33% in the concentrations of H_2O_2, O_3, and NO_y.

The dependence of H_2O_2 concentration on O_3 and NO_y is very different in different regions of this plume. There is an inverse relationship between the concentrations of H_2O_2 and O_3 ($r^2 = 0.76$) and between H_2O_2 and NO_y ($r^2 = 0.86$) observed during the 630-m transect (region B in Figure 7), as on August 27th. However, on the 1240-m transect (region C, Figure 7), there is a positive correlation between H_2O_2 and O_3 ($r^2 = 0.89$)

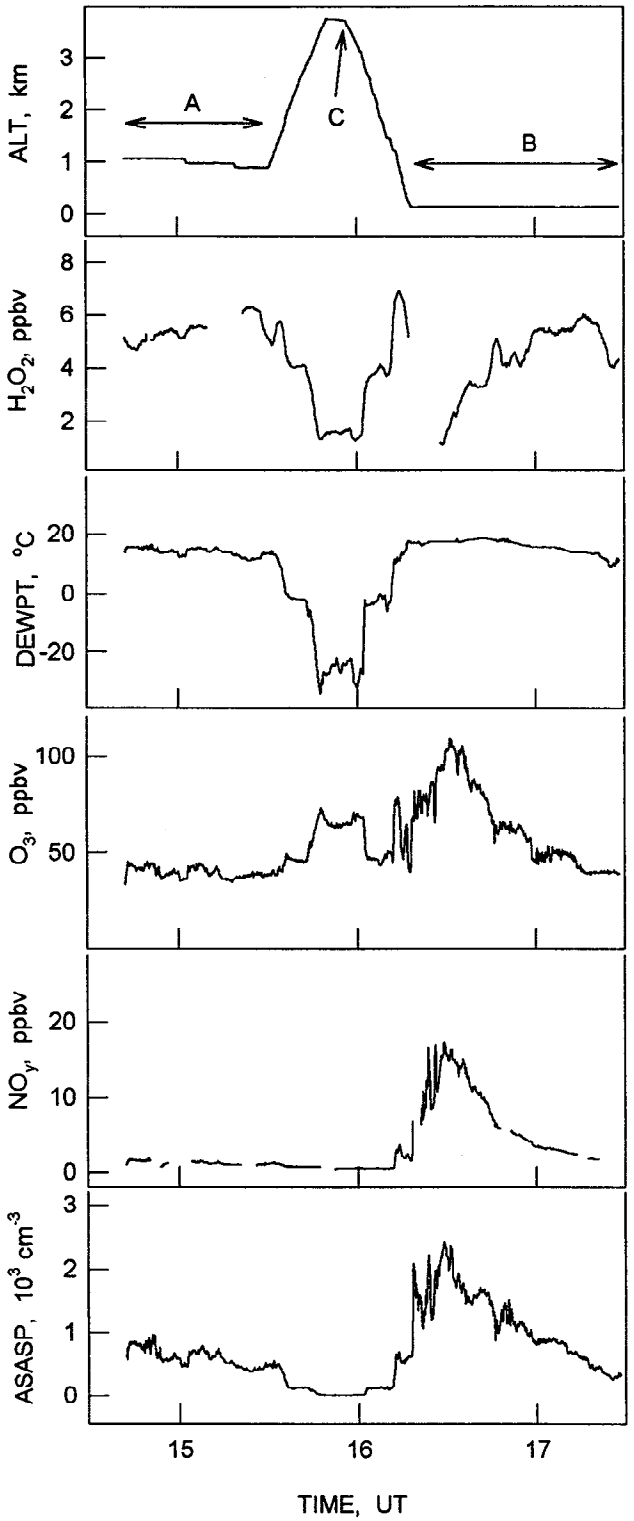

Figure 4. Concentrations of selected species on August 27 as a function of time. The horizontal transect indicated as region A and the plume encountered in the region shown as B are described in the text.

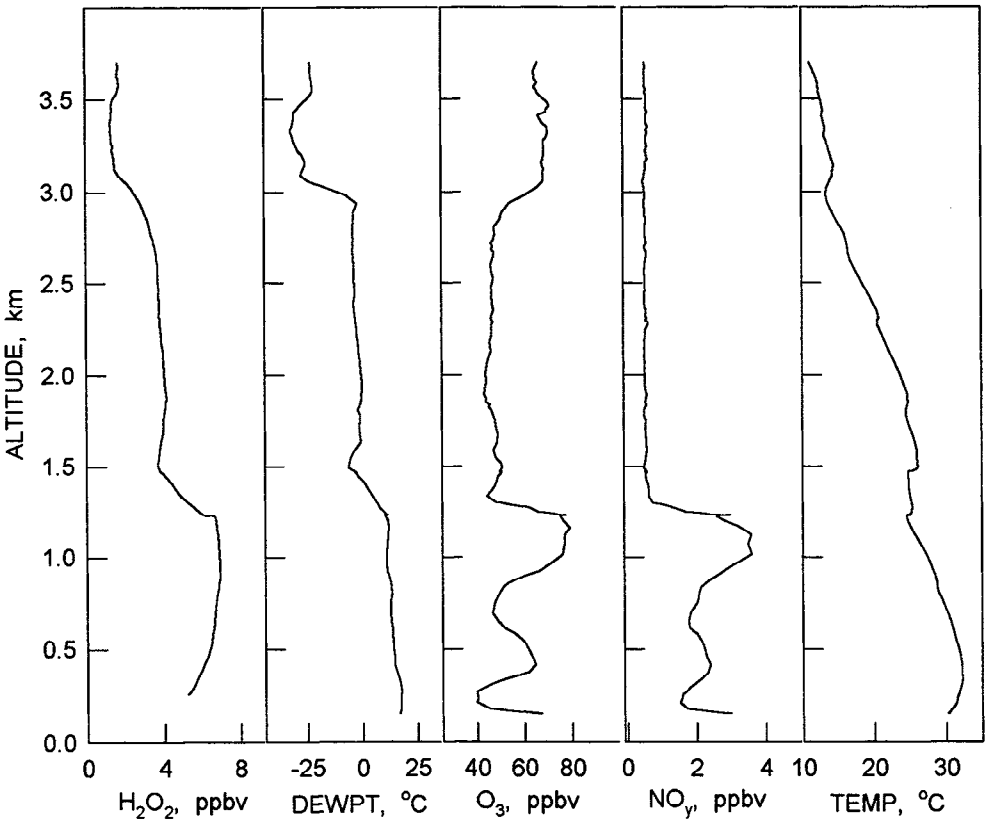

Figure 5. Vertical distribution of selected species on August 27 illustrating pollutant layer traversed during sounding at point C in Figure 4.

and between H_2O_2 and NO_y ($r^2 = 0.90$). In contrast to the plume sampled on August 27, this plume is estimated to have taken ~72 hours to reach Nova Scotia from the source region, consistent with the observation that virtually all of the NO_x had been depleted.

August 31. Elevated concentrations of H_2O_2, O_3, and NO_y were again observed on the August 31 flight. The first portion of the flight, passing over land to the coast of Maine at an altitude of ~1 km, exhibited fairly low concentrations of these species (region A in Figure 9). On the southward leg to 42.5°N, 68°W (region B in Figure 9), the concentration of H_2O_2 rose to ~8, O_3 to ~70, and NO_y to ~4 ppbv. A sounding at this location (point C in Figure 9) revealed the presence of a layer containing high concentrations of NO_y and aerosol particles at ~1500 m. The vertical profile for this sounding, illustrated in Figure 10, shows the layers to be somewhat more diffuse than on earlier days, due perhaps to the absence of a well-defined inversion above the surface. The high-NO_y air mass sampled during the return leg, region D in Figure 9, is well aged, with virtually no NO_x remaining, as shown by the lower trace in the NO_y panel in Figure 9. In contrast to the well-aged plumes observed on August 27 and 28, this air mass shows no correlation between H_2O_2 and NO_y or between H_2O_2 and O_3.

Peroxide Relationships in the Free Troposphere

In the free troposphere, where NO_x concentrations are low and peroxides are the principal radical sink, peroxide formation is expected to depend on ozone photolysis rate and the concentrations of ozone and water vapor [*Kleinman*, 1986; 1991]. Figure 11 shows that the concentration of H_2O_2 correlates strongly ($r^2 = 0.80$) with the product of concentrations of H_2O and O_3 for altitudes above 1500 m. Over 90% of the points in this high-altitude subset exhibit NO_y concentrations below 1.5 ppbv, and the correlation is lost when low altitude, higher NO_y data is included.

The regression line in Figure 11 is what we expect based on a simple steady state model in which the concentration of H_2O_2 is linearly related to the production rate of free radicals, which in turn is approximately proportional to the product of water vapor and ozone. Figure 11 indicates that this model can account for most of the variance in the low NO_y data subset. However, a change in the relation between H_2O_2 and O_3*H_2O is evident at low radical production rates, that is, values of O_3*H_2O below 100 ppm^2. This may be due to the dependence of H_2O_2 lifetime on OH concentration, together with the dependence of OH concentration on radical production rate. Detailed photochemical model calculations are needed to investigate this possibility.

Correlation Between O_3 and (NO_z + $2H_2O_2$)

The relationship between H_2O_2, O_3 and NO_y for plume data can be understood in terms of a correlation noted by *Sillman* [1995] in his model calculations of O_3-NO_x-ROG sensitivity. Sillman observed that O_3 concentration correlated with the sum of H_2O_2 and NO_x oxidation products, that is, NO_y-NO_x or NO_z; the correlation was strongest over water where model deposition rates were low. This relationship emphasizes O_3 as radical source, and the quantity NO_z + $2H_2O_2$ as radical sink, allowing for the 2:1 stoichiometry in forming H_2O_2 from HO_2 radicals. It applies whether the dominant radical sink is peroxide or HNO_3 formation, and whether the air mass is fresh or aged.

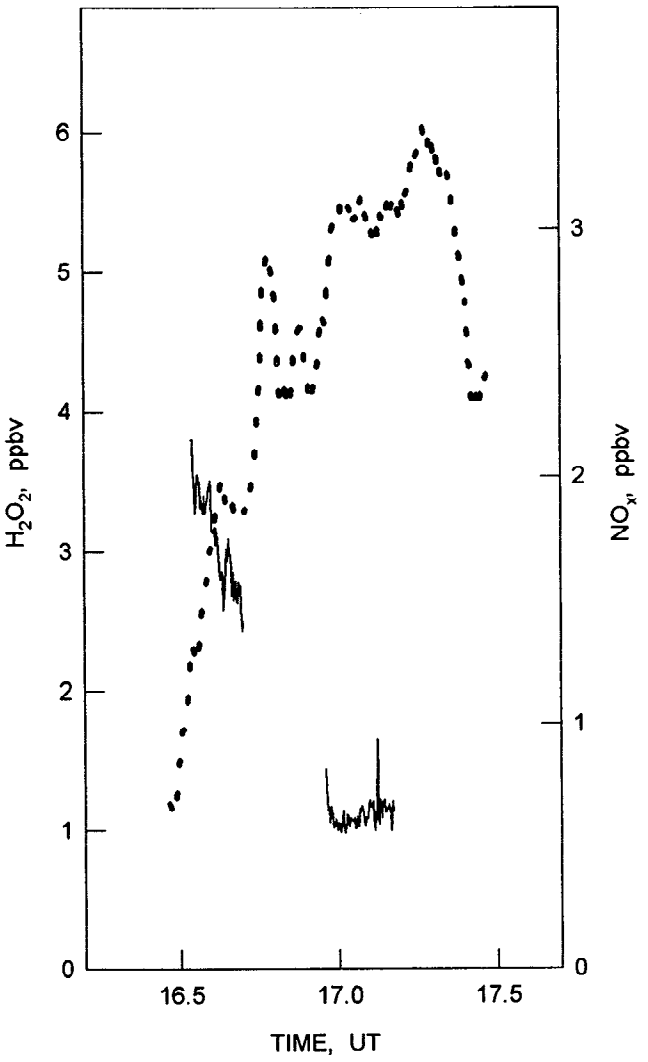

Figure 6. Inverse relation between H_2O_2 (dotted line) and NO_y (solid line) on August 27 during a plume transect at 145 m at 16.4 UT.

Although individual subsets of the NARE data exhibit very different relationships among H_2O_2, O_3, and NO_y, as described in the case studies above, the complete data set exhibits a linear correlation (r^2 = 0.73) between O_3 and NO_z + $2H_2O_2$, as illustrated by *Daum et al.* [this issue]. The lack of vertical mixing and wet deposition prevailing during this period facilitated the observation of this relationship.

Summary

Peroxide concentrations varying from the detection limit to over 11 ppbv were observed over the eastern North Atlantic in the summer of 1993, with the highest concentrations occurring in well-defined layers above the surface inversion. The composite data set exhibits only a shallow concentration maximum near the top of the boundary layer (~2 km), in contrast to other aircraft studies conducted over the continent where the boundary layer is well mixed [*Heikes et al.*, 1987; *Boatman et al.*, 1990; *Tremmel et al.*, 1993]. Plumes of high H_2O_2 concentration frequently were found in well-aged air masses advected from the continent along with high concentrations of anthropogenic pollutants. The observation of high H_2O_2 concentrations above the surface inversion underscores the difficulty in interpreting surface measurements of photochemically important species.

Total peroxide was dominated by H_2O_2, with MHP concentrations near 0.5 ppbv observed on only 3 of 14 flights. The absence of MHP is interesting in light of the high H_2O_2 concentrations observed in plumes, and merits further investigation.

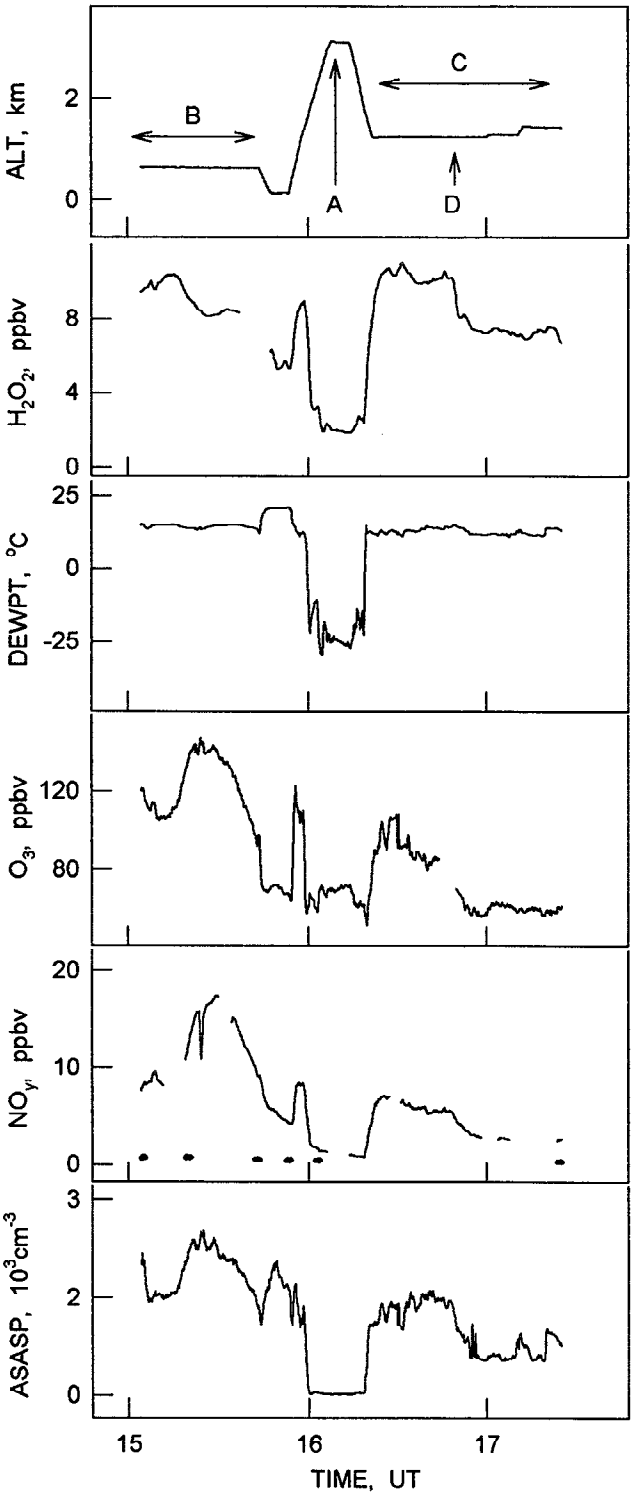

Figure 7. Concentrations of selected species on August 28 as a function of time. Horizontal transects, indicated by regions B and C are described in the text. Dashed line in the NO_y panel represents NO_x.

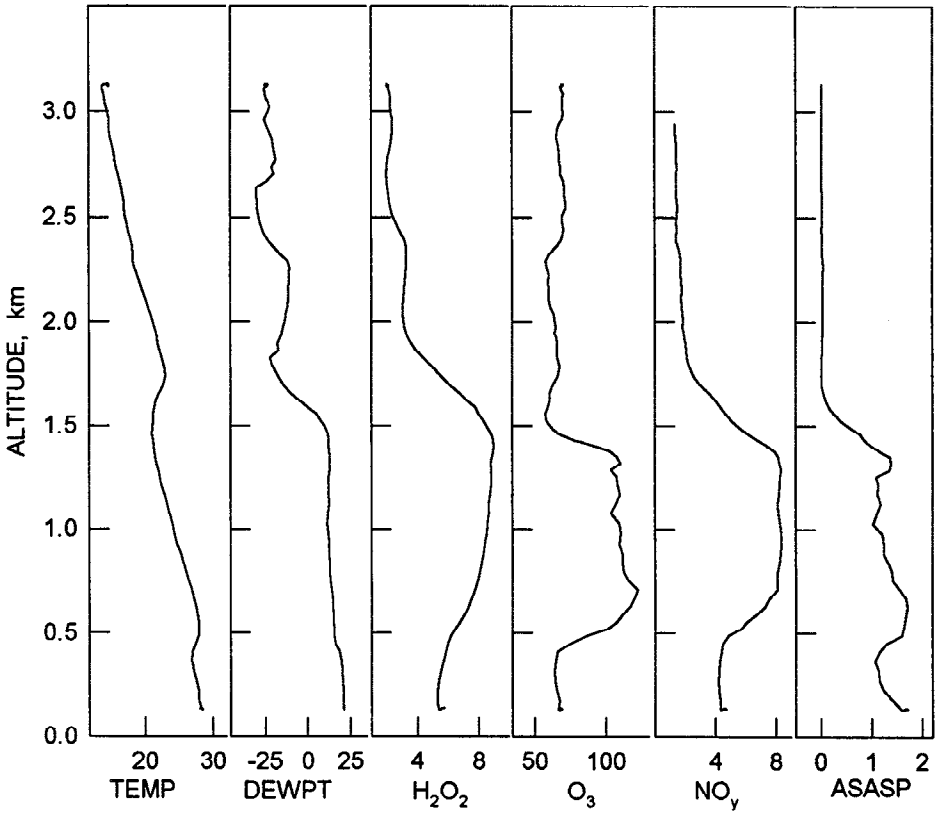

Figure 8. Vertical distribution of selected species at August 28 illustrating pollutant layer traversed during sounding at point A in Figure 7. Units are the same as in Figure 7.

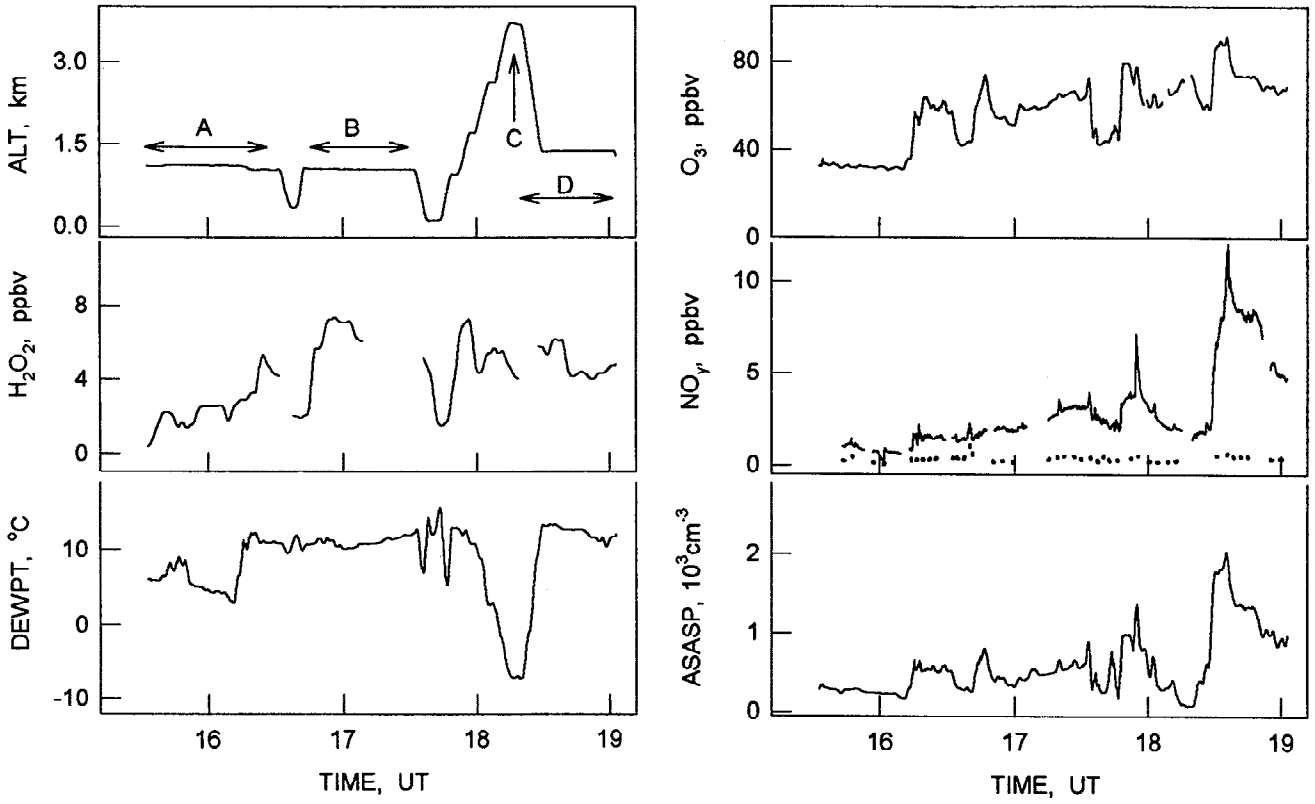

Figure 9. Concentrations of selected species on August 31 as a function of time. Horizontal transects, indicated by regions A, B, and D are described in the text. Dashed line in the NO_y panel represents NO_x.

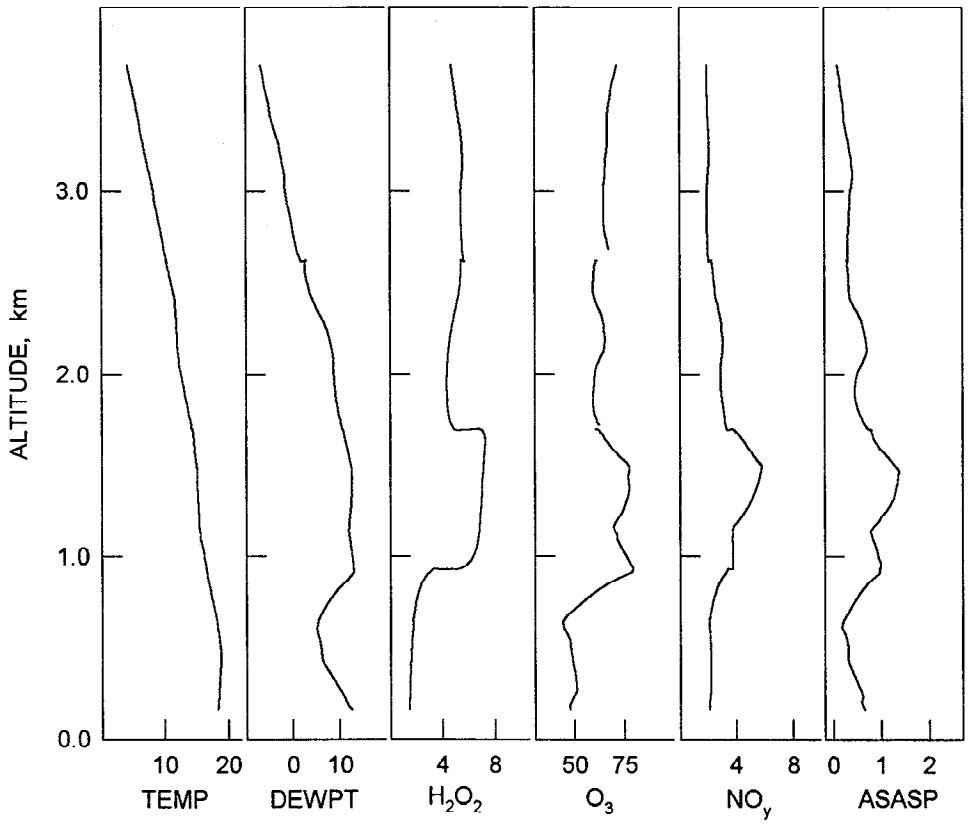

Figure 10. Vertical distribution of selected species on August 31 illustrating pollutant layer traversed during sounding at point C in Figure 11. Units are the same as in Figure 9.

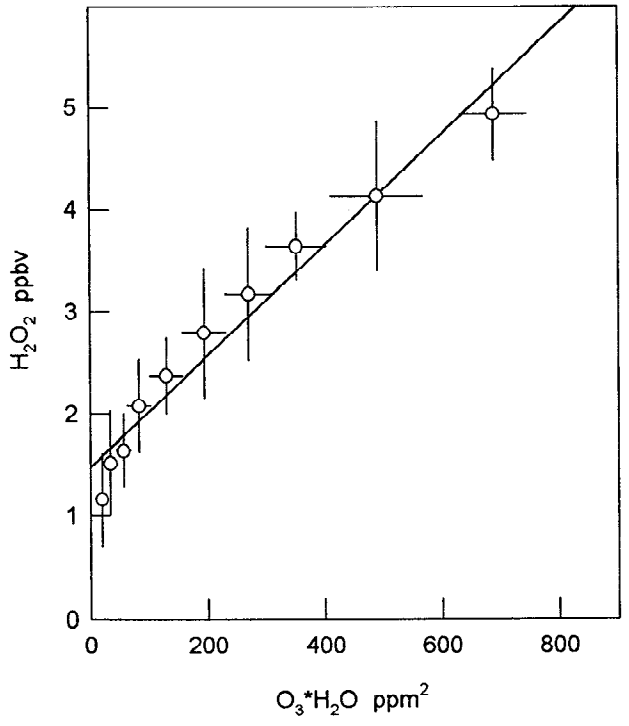

Figure 11. Correlation between H_2O_2 and the product of O_3 and H_2O in the free troposphere, i.e. data with $[NO_y]<1.5$ ppbv. Data were sorted in order of increasing x value, and collected in 10 groups of equal size; the mean is indicated by a circle and lines show the range of x and y values for each group. A linear regression through the means is included.

A strong correlation between H_2O_2 concentration and the product H_2O*O_3 was observed in "clean" free tropospheric air, that is, air masses with NO_y concentrations below 1.5 ppbv, confirming our understanding of the formation mechanism for H_2O_2 under conditions of low NO_x and minimal depositional losses.

In pollutant plumes, the relationship between H_2O_2 concentration and that of other species is complicated. These regions are characterized by high concentrations of NO_y, O_3 and aerosol particles, indicating that they have been subjected to substantial photochemistry at the source region and during transport to Nova Scotia. Peroxide concentrations are also high in these plumes. Individual horizontal transects within polluted layers exhibit H_2O_2 concentration with the inverse dependence on O_3, and NO_x characteristic of fresh air masses, or the direct relationship expected in NO_x-depleted air. Detailed model calculations are required to unravel these different behaviors.

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